IN THE CLAIMS

This listing of claims replaces all prior versions, and listings, in this application.

1. (currently amended) A drug, containing:

metabolic product-biomass prepared by incubating *Rhodopseudomonas*capsulatas FERMBP-7434 strain a photosynthetic bacterium together with a lactic acid bacterium so as to cause the photosynthetic bacterium to produce a biomass comprising a viscous material;[[,]] wherein after being subjected to water-washing and subsequently to acid hydrolysis, the biomass has a glucose content ranging from 0.8 to 3.3 weight %, a ribose content ranging from 0.2 to 1.0 weight %, a rhamnose content ranging from 0.4 to 2.0 weight %, and a fucose content of 0.6 weight % or less

the photosynthetic bacterium being Rhodopseudomonas capsulatas FERMBP-7434 strain.

2. (currently amended) The drug as set forth in Claim 1, wherein:

the <u>biomass metabolic product</u> contains <u>bacteriochlorophyll-bacteriocholorophyll</u> in a range of from 0.2 to 3.0 weight % (weight %).

- 3. (currently amended) The drug as set forth in Claim 1, wherein:
- the <u>biomass</u> metabolic product contains <u>bacteriochlorophyll</u> bacteriocholorophyll in a range of from 0.6 to 1.9 weight % (weight %).
- 4. (currently amended) The drug as set forth in Claim 1, wherein:

the <u>biomass</u> metabolic product contains a carotinoid material in a range of 0.5 to $7.5 \, \mu \text{mol/g} \cdot (\mu \text{mol/g})$.

5. (currently amended) The drug as set forth in Claim 1, wherein:

the <u>biomass</u>-metabolic product contains a carotinoid material in a range of 2.4 to μ mol/g-(μ mol/g).

6. (currently amended) The drug as set forth in Claim 1, wherein:

after being subjected to acid hydrolysis, the biomass metabolic product has a glucose content contents (weight %) ranging from 2.4 to 7.5 weight %, a ribose content contents (weight %) ranging from 0.3 to 1.1 weight %, a rhamnose content contents (weight %) ranging from 1.0 to 3.3 weight %, and a fucose content contents (weight %) ranging from 0.6 to 2.6 weight %.

7. (currently amended) The drug as set forth in Claim 1, wherein:

after being subjected to acid hydrolysis, the biomass metabolic product has a glucose content contents (weight %) ranging from 3.5 to 6.5 weight %, a ribose content contents (weight %) ranging from 0.4 to 1.0 weight %, a rhamnose content contents (weight %) ranging from 1.2 to 3.0 weight %, and a fucose content contents (weight %) ranging from 0.8 to 2.4 weight %.

Claim 8 (canceled) The drug as set forth in Claim 1, wherein:

9. (currently amended) The drug as set forth in Claim 1, wherein:

after being subjected to water-washing and subsequently to acid hydrolysis, the biomass-metabolic product has a glucose content contents (weight %) ranging from 1.0 to 3.0 weight %, a ribose content contents (weight %) ranging from 0.3 to 0.9 weight %, a rhamnose content contents (weight %) ranging from 0.5 to 1.6 weight %, and a fucose content-contents (weight %) of 0.5 weight % or less.

Claims 10-13 (cancel)

- 14. (currently amended) The drug as set forth in Claim 1, wherein: the lactic acid bacterium is <u>Lactobacillus</u> Lactobacillus spp.
- 15. (currently amended) The drug as set forth in Claim 1, wherein:
 the lactic acid bacterium is <u>Lactobacillus bulgalicus</u> Lactobacillus bulgalicus.

16. (currently amended) A method of manufacturing a drug, comprising the steps of:
incubating in a liquid medium Rhodopseudomonas capsulatas FERMBP-7434

strain a photosynthetic bacterium together with a lactic acid bacterium-so as to cause
the photosynthetic bacterium to produce a biomass comprising a viscous material in the
[[a]] liquid medium, the photosynthetic bacterium being Rhodopseudomonas capsulatas
FERMBP-7434-strain; and

separating the biomass comprising a viscous material a metabolic product from the liquid medium.